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CLAIMS

1. A method of preparation of (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide of formula I

$$H_2NO_2S$$
 O
 NH_2
 I

10 characterized in that

a. a protecting group is introduced to N-[(1R)-2-(4-methoxyphenyl)-1-methylethyl]-N-[(1R)-1-phenylethyl)]amine of formula VIII

to obtain an amide of formula IX

wherein A can be an acyl having 2 to 8 carbons,

b. whereupon the amide of formula IX is chlorosulfonated and the resulting sulfochloride is converted to a sulfonamide of formula X

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wherein A is as defined above,

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- c. and the sulfonamide of formula X is hydrogenated to obtain the compound of formula I.
- 2. The method according to claim 1 c h a r a c t e r i z e d in that the protecting group A is an acyl, preferably acetyl.
- 3. The method according to claim 2 c h a r a c t e r i z e d in that acetanhydride at 50 to 100 °C is used as the acetylation agent.
 - 4. The method according to claim 1 c h a r a c t e r i z e d in that the sulfochloride resulting from chlorosulfonation is not isolated and is directly converted to the sulfonamide with ammonia.
- 5. The method according to claim 4 c h a r a c t e r i z e d in that chlorosulfonation takes place in methylenechloride at -30 to +30 °C.
 - 6. The method according to claim 1 c h a r a c t e r i z e d in that hydrogenation is carried out under catalysis with palladium.
 - 7. The method according to claim 6 c h a r a c t e r i z e d in that the catalyst is 3% Pd/C with 50% water content at a pressure of 1 to 5 MPa and a temperature of 50 to 100 °C.
 - 8. A method of preparation of (R)-(-)-5-[2-[2-(2-ethoxyphenoxy)ethylamino]propyl]-2-methoxybenzenesulfonamide of formula II

II

c h a r a c t e r i z e d in that the intermediate of formula I produced according to any of the preceding claims is used for the synthesis.

9. The method according to claim 8 c h a r a c t e r i z e d in that intermediate I is reacted with a compound of formula IV

10. A sulfonamide of formula X

$$H_2NO_2S$$
 O
 X
 X

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wherein A is as defined in claim 1.

11. The sulfonamide according to claim 10, wherein A is acetyl.